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	Practical challenges
	 Legal and regulatory hurdles e.g. Medicaid's best price rule, Off-label use, anti-kickback statute, data privacy issues
	 Contractual or financial flow issues Payer who agrees the price with the manufacturer may be reimbursing the provider who in turn pays the wholesaler who pays the manufacturer
	 Data collection that tracks uses and outcomes by indication Proxies or surrogate measures: e.g. treatment duration?
	Arbitrage (re-selling) must be difficult
	 How to attribute value between drugs for combination therapies?
Office of Health	4. Can innovative payment models work?

	Conclusion	\rightarrow
	Short term rewards of greater patient a term gains of incentivising R&D and cor	ccess, long npetition
	 In the short term, IBP can improve overall well access increases, but expenditure may rise 	fare if patient
	 Existing research has neglected longer term in optimised incentives for R&D can lead to new to options for patients 	npact: treatments
	 Increased price competition at the indication-le down prices and <i>delivers better value to the he</i> The UK NHSE competitive tendering process for drugs separates tenders by genotype – in effect US health plans and PBMs are currently piloting a with the objective to better manage expenditure 	evel drives ealth system Hepatitis C by indication IBP approaches
)ffice of lealth conomics Research		5. Conclusion

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